

Fluorescence Signal

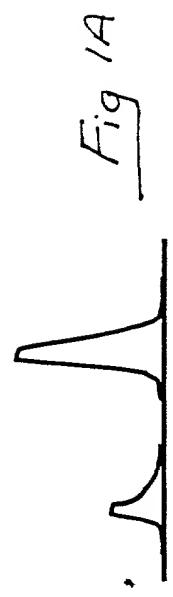


Fig 1A

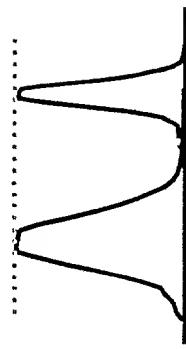
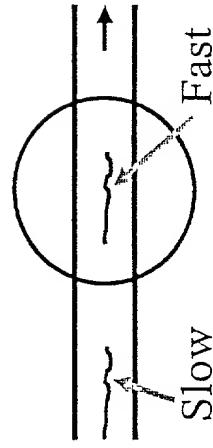
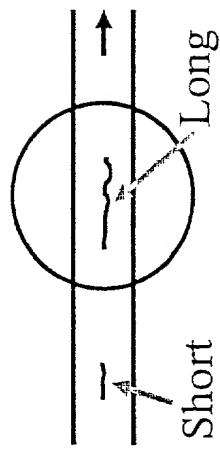
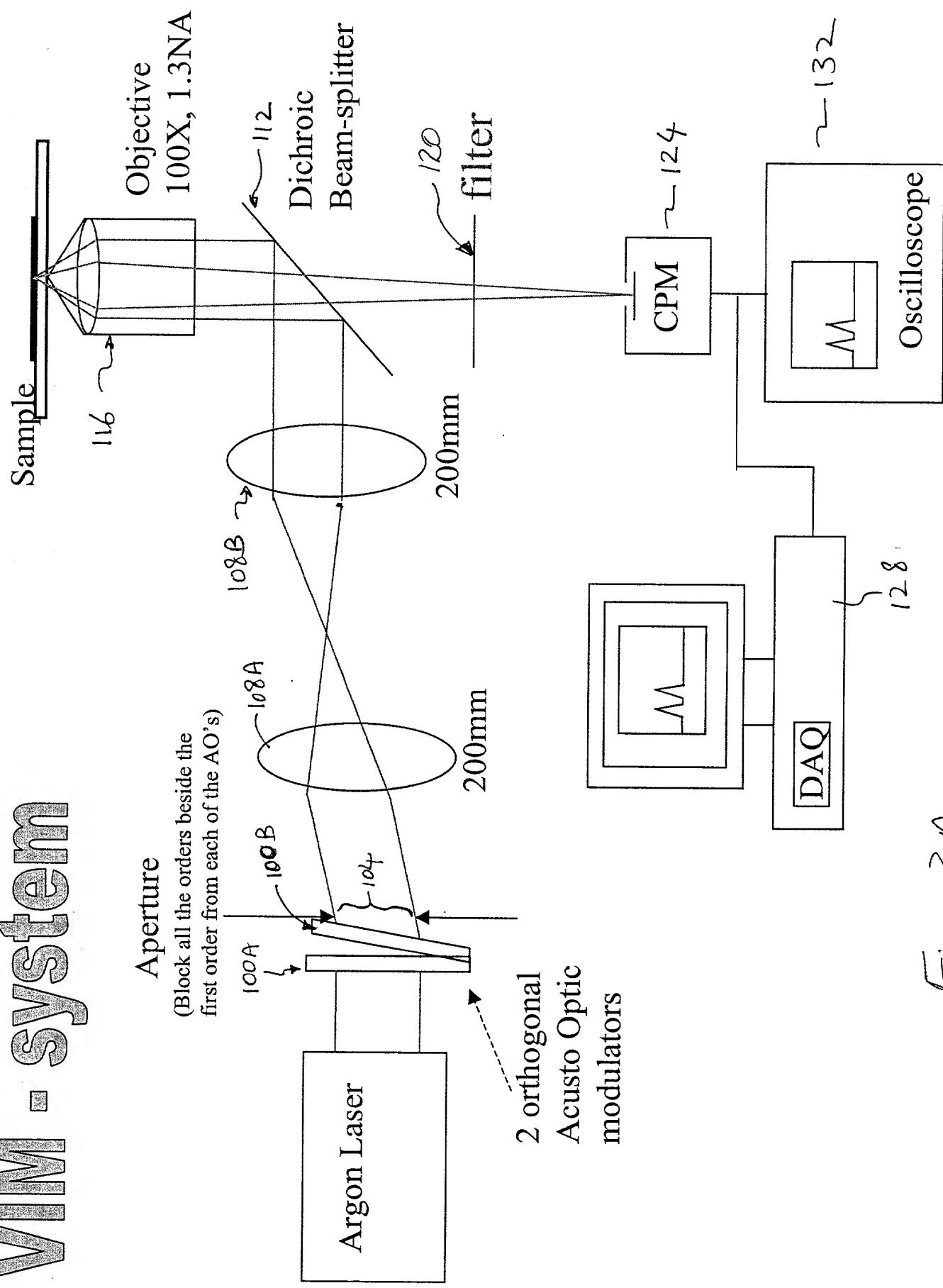
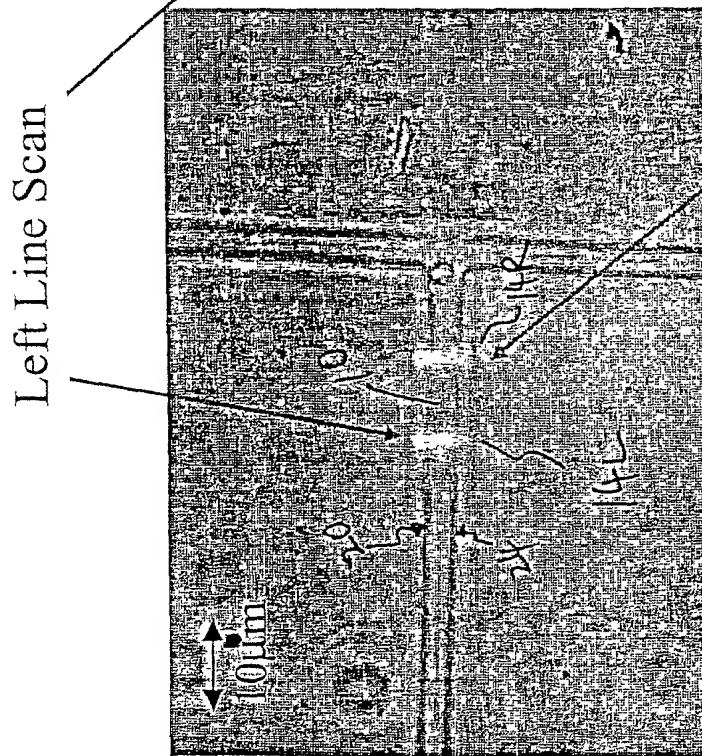
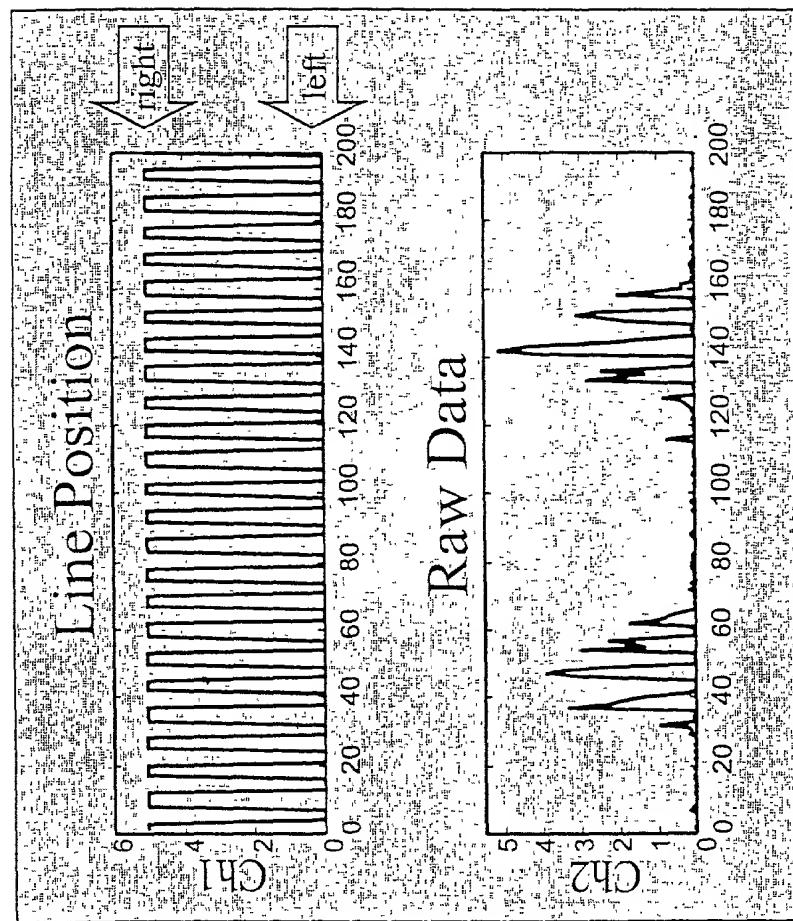


Fig 1B



VIM - System





The beam after the two Acusto Optics Modulators

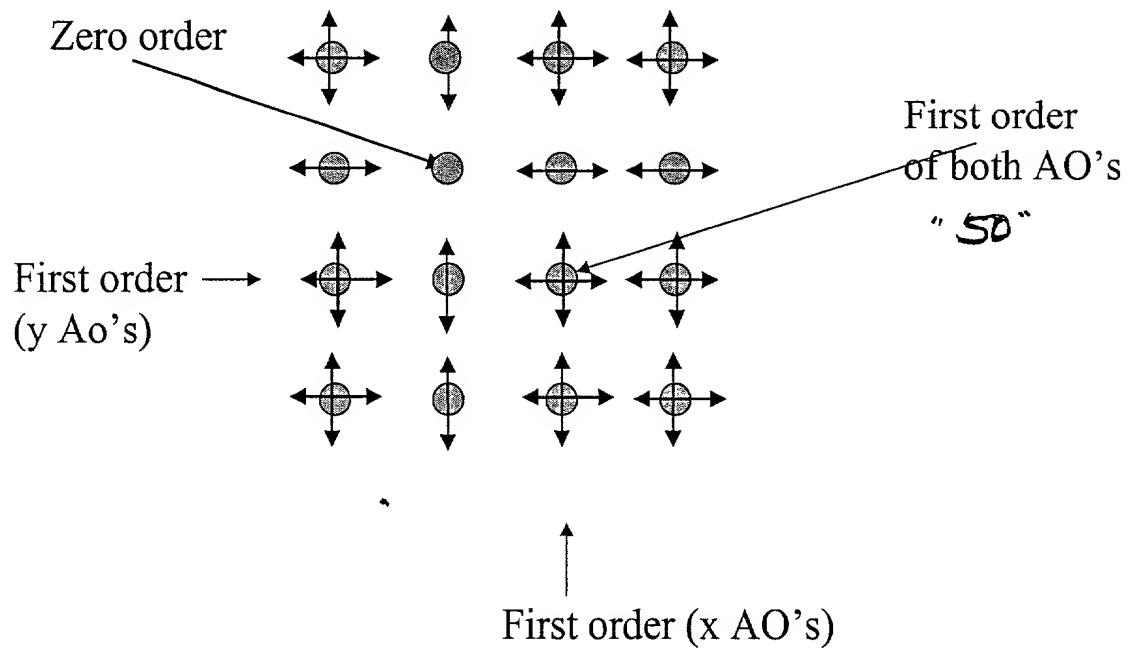
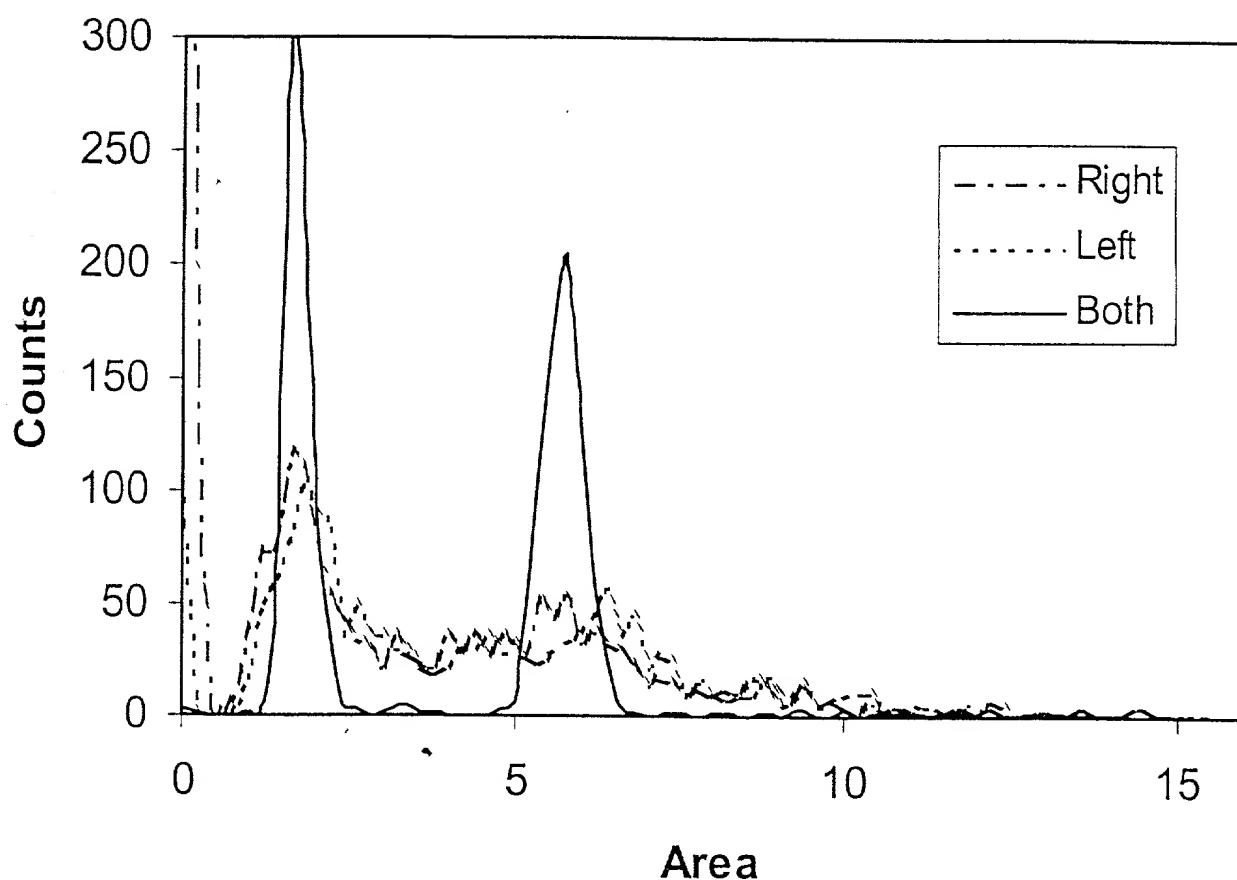
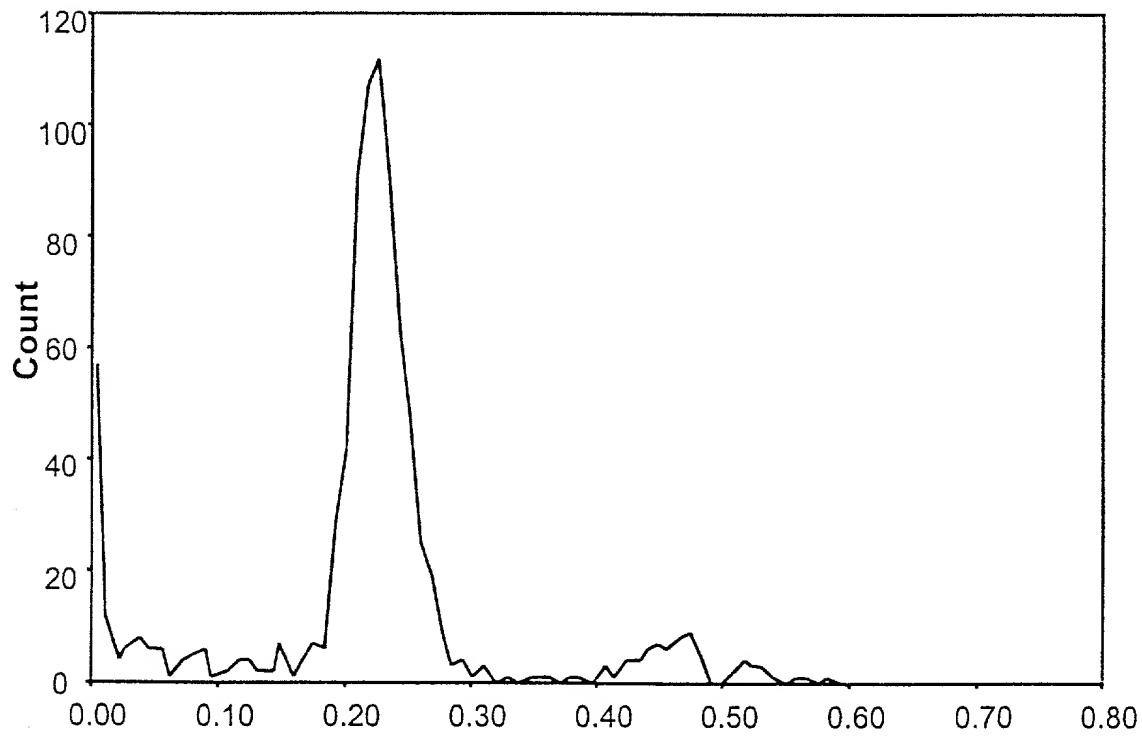


Fig 2c



Area

Fig 3



Area

Fig 6

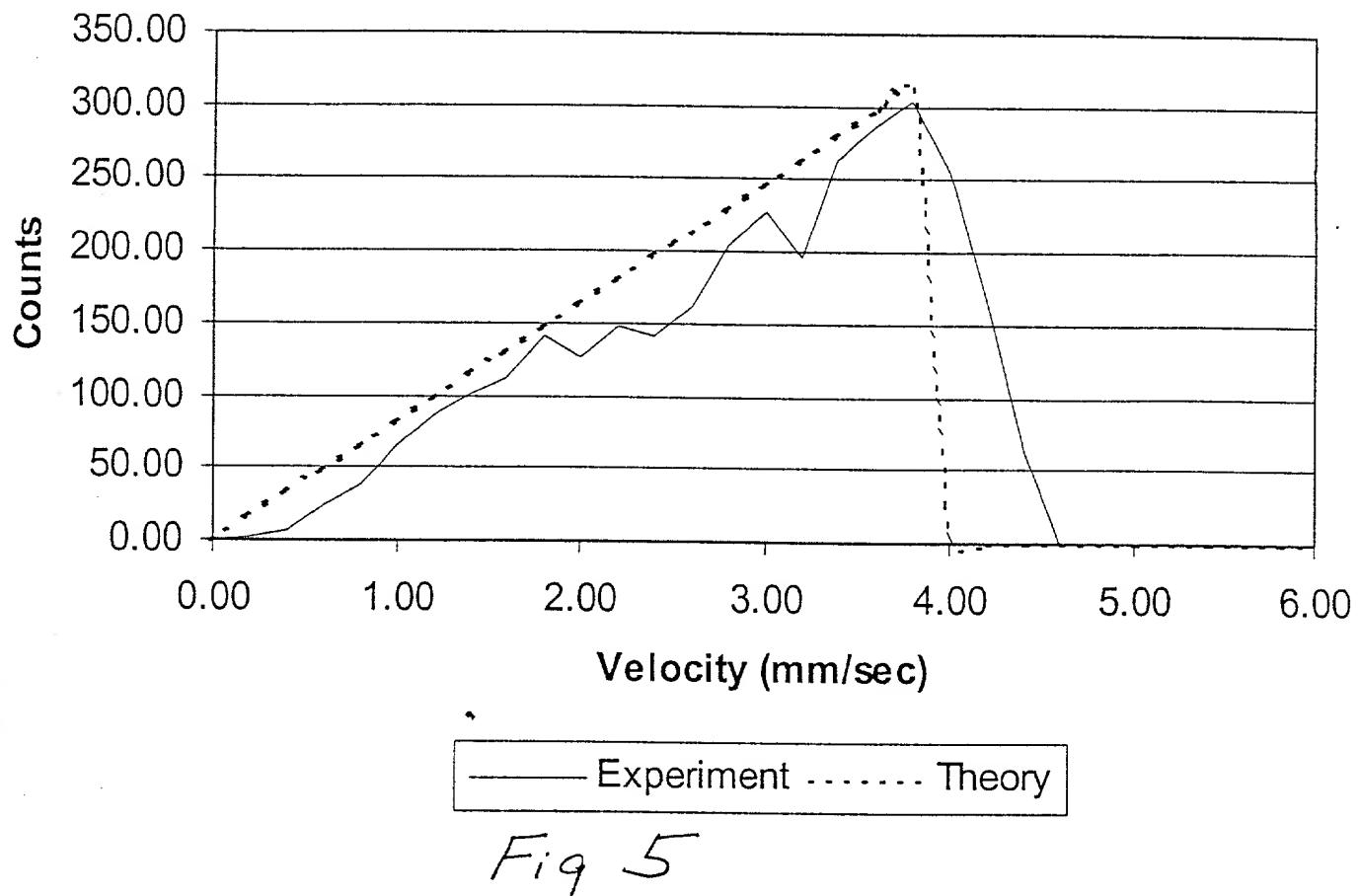
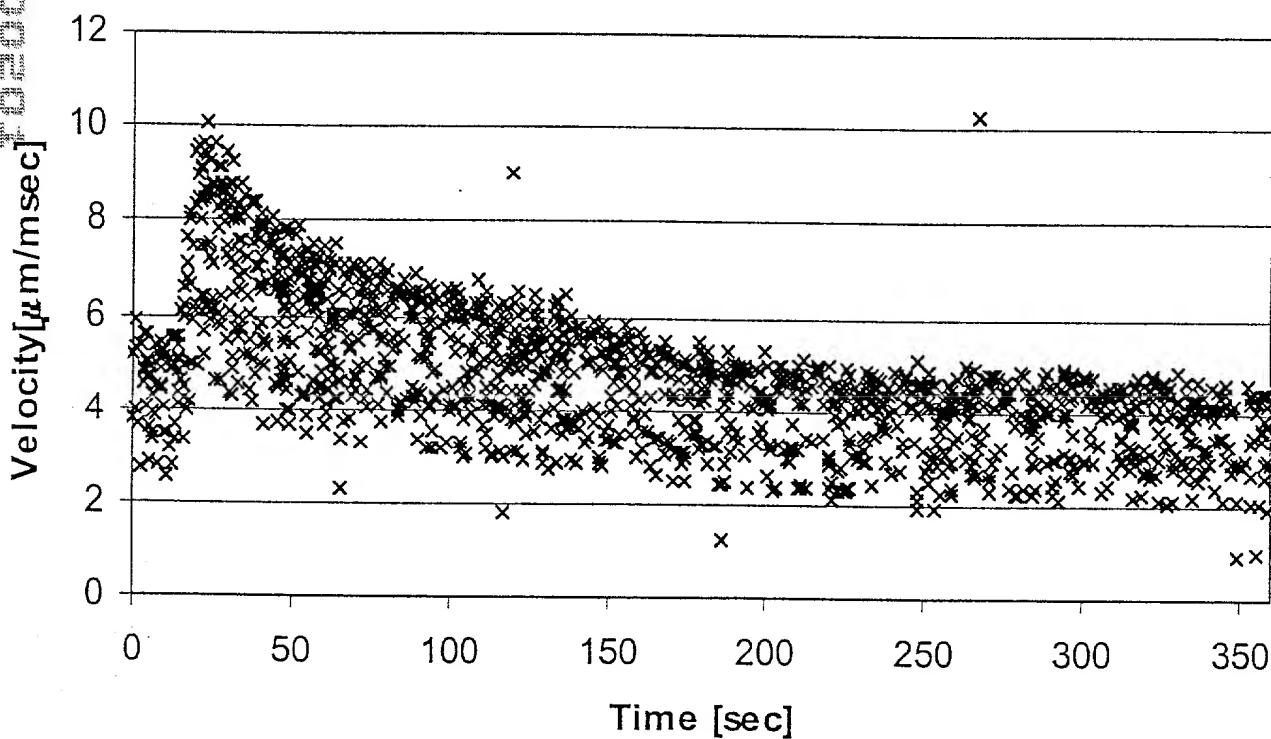


Fig 1 Fig 2 Fig 3 Fig 4 Fig 5 Fig 6 Fig 7



ChDiv

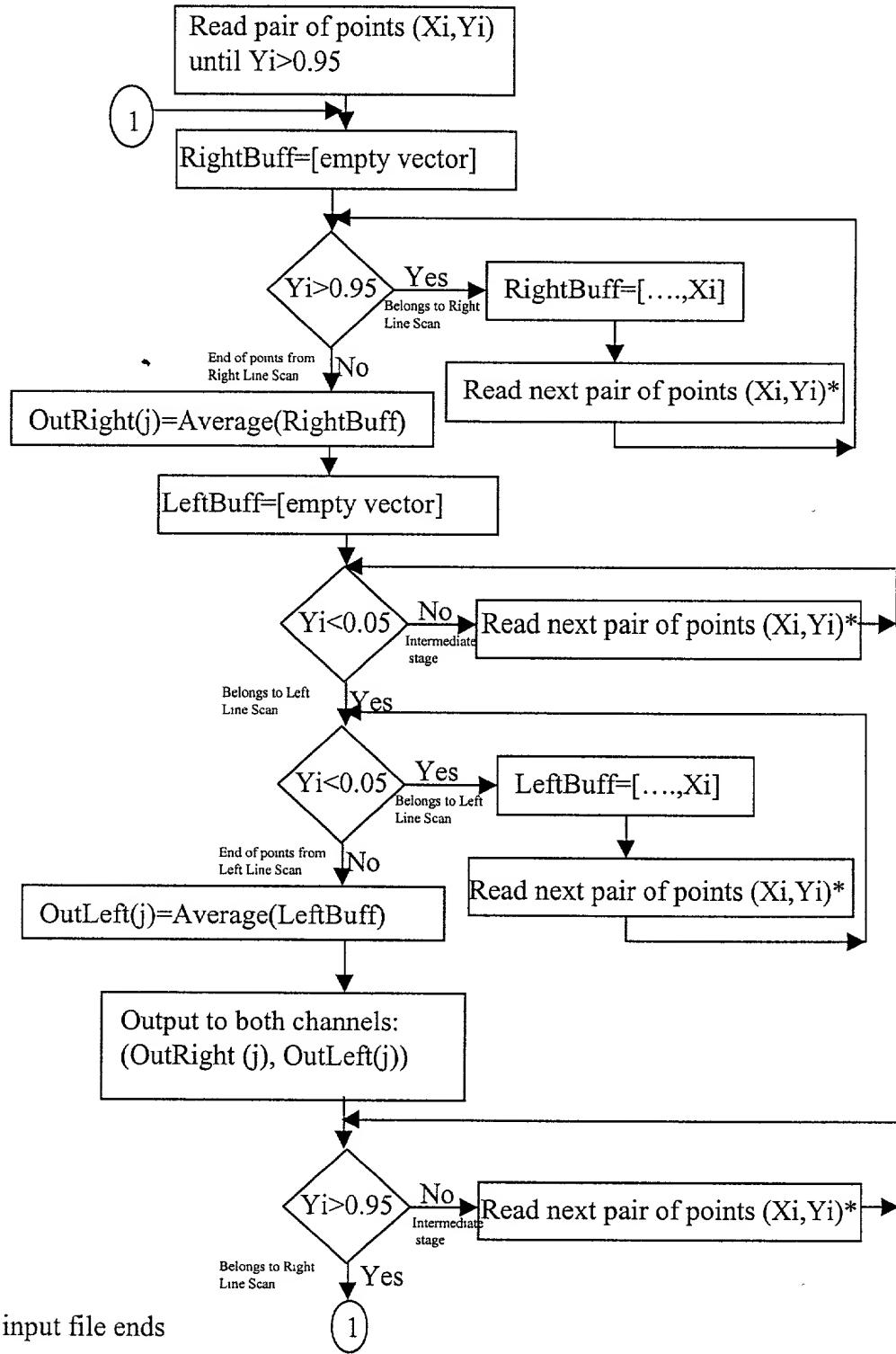
Input - two vectors: $Y(i)$ - channel 1 - square wave - chopping signal, $0 \leq Y(i) \leq 1$
 $X(i)$ - channel 2 - fluorescence raw data - from the detecting region (both line scan)

Usually Sampled
at 40KHz

Output - two vectors: $OutRight(j)$ - fluorescence from Right Line scan
 $OutLeft(j)$ - fluorescence from Left Line scan

Usually Sampled
at 5KHz

The sampling rate of the output channels always equals the frequency of the chopping signal



* Program ends when input file ends

Fig 7

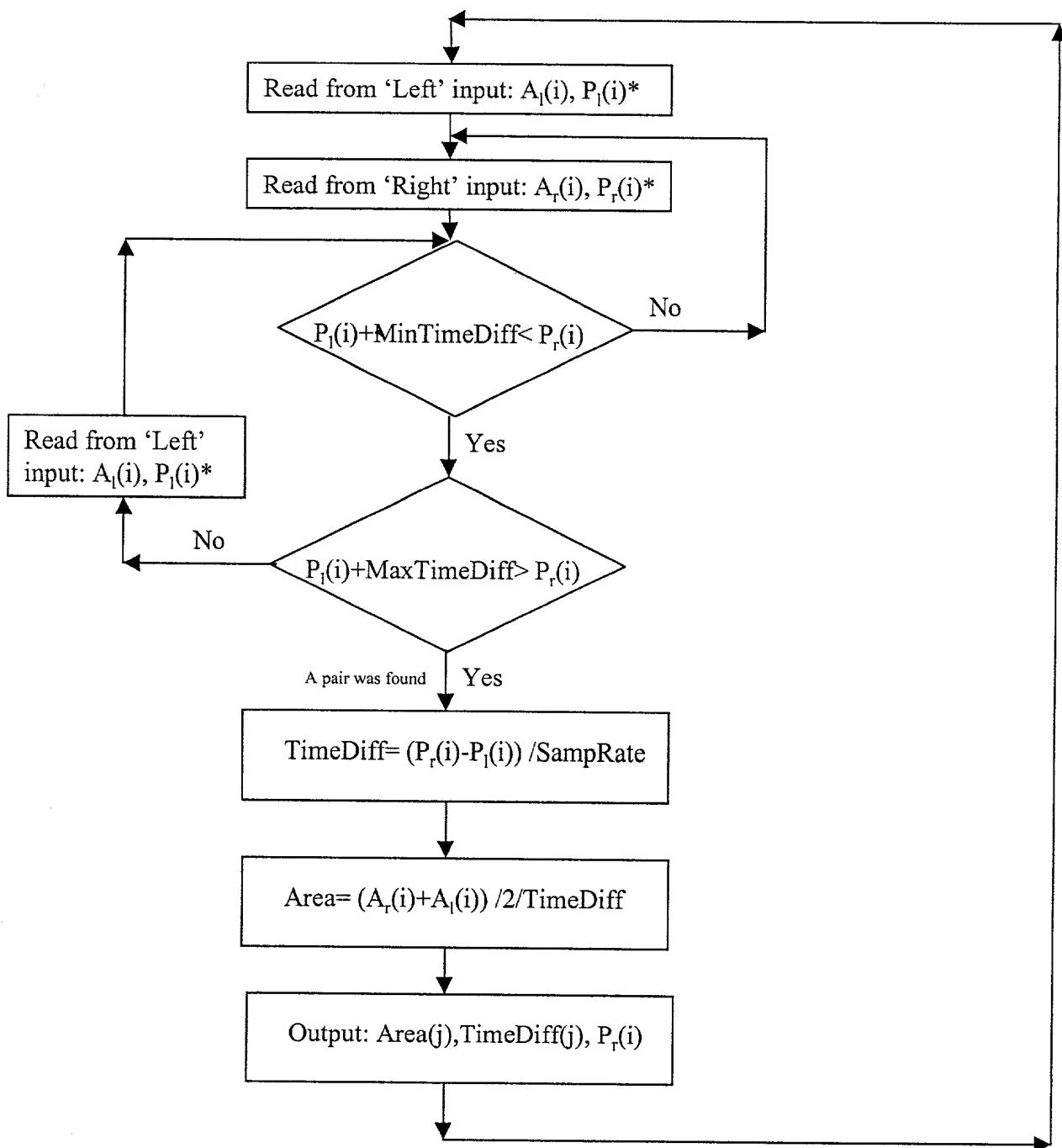
ArVIAnalyzer

Input: two files (one for each line scan).

Each file contain 2 vectors one of Positions ($P_l(i)$) and the other has the corresponding Area ($A_l(i)$)

Output: three vectors - Area, TimeDiff (inversely proportional to velocity), Position

Parameters that can be determined - MinTimeDiff, MaxTimeDiff



Position is presented in point number and not time

TimeDiff is in Seconds and is inversely proportional to the velocity

* Program ends when one of the input files ends

Fig 8